```
1978-42505A [24] WPIX
AN
     Pharmaceutical compsn. containing self-emulsifying oil e.g. silicone - for
TI
     admin. as capsules, giving rapid gastrointestinal resorption
DC
    A96; B07
    GROVES M J
TN
     (SCHB-C) SCHERER CORP R P
PA
CYC 4
     DE 2753526 A 19780608 (197824)* DE JP 53072814 A 19780628 (197831) JA FR 2372635 A 19780804 (197836) FR IT 1090703 B 19850626 (198634) IT
                                                                                <---
PI
PRAI GB 1976-50577
                          19761203
     IC A61K047-00; A61K009-08
IPCR A61K0047-06 [I,A]; A61K0047-06 [I,C]; A61K0047-14 [I,A]; A61K0047-14
     [I,C]; A61K0047-24 [I,A]; A61K0047-24 [I,C]; A61K0009-48 [I,A];
     A61K0009-48 [I,C]
EPC A61K0009-48H4
                     UPAB: 20050417
     DE 2753526 A
AB
     Pharmaceutical compsn. contains a unit dose of the active ingredient (I)
     and a self-emulsifying oil (II). Suitable (II) are steroids; antimycotics;
     tranquilisers, sedatives; diuretics; anti-inflammatories; analgesics;
     antibiotics and insulin; specifically dexamethasone, prednisolone,
     griseofulvin, diazepam, chlordiazepoxide, furosemide, indomethacin,
     lipophilic penicillins and bleomycin.
     (I) and (II) are presented in solution or suspension, and (II) consists of
     oil (IIa), e.g. a silicone or triglyceride, and a surfactant (IIb), pref.
     nonionic e.g. polyethoxylated alkylphenols.
     For rectal, vaginal and especially oral administration; the compsns. can be
     filled into capsules to give highly reproducible doses, and when ingested
     have improved resorption from the gastro-intestinal tract. CPI: A12-V01; B04-B01B; B04-B01C; B04-C03; B12-A02; B12-C08; B12-C10;
MC
           B12-D01; B12-D07; B12-G03; B12-G04; B12-M09; B12-M10
    ANSWER 2 OF 2 HCAPLUS COPYRIGHT 2009 ACS on STN
T.1
    1978:465260 HCAPLUS
     89:65260
DN
OREF 89:10009a,10012a
   Entered STN: 12 May 1984
    Pharmaceutical preparation and administration form
TN
     Groves, Michael John
    R. P. Scherer Ltd., UK
PA
   Ger. Offen., 22 pp.
    CODEN: GWXXBX
DT
     Patent
LA
    German
    A61K009-08
    63-5 (Pharmaceuticals)
CC
FAN.CNT 1

    KIND
    DATE
    APPLICATION NO.
    DATE

    A1
    19780608
    DE 1977-2753526
    19771201

    A
    19790607
    AU 1977-31116
    19771201

    PATENT NO.
     _____
                                                                       19771201 <--
PT
     DE 2753526
                          A 19790607
A 19780628
     AU 7731116
                                              JP 1977-144921
                                                                      19771202
     JP 53072814
                          A1 19780630
                                             FR 1977-36418
     FR 2372635
PRAI GB 1976-50577
                          A
                                19761203
CLASS
PATENT NO. CLASS PATENT FAMILY CLASSIFICATION CODES
 IC
DE 2753526
                         A61K009-08
                 TPCT
                         A61K0009-08 [ICM]
                 TPCR
                         A61K0047-14 [I,C*]; A61K0047-14 [I,A]; A61K0009-48
                         [I,C*]; A61K0009-48 [I,A]; A61K0047-06 [I,C*];
                         A61K0047-06 [I,A]; A61K0047-24 [I,C*]; A61K0047-24
                         [I,A]
```

```
A61K0047-14 [I,C*]; A61K0047-14 [I,A]; A61K0009-48
                  TPCR
                         [I,C*]; A61K0009-48 [I,A]; A61K0047-06 [I,C*];
                         A61K0047-06 [I,A]; A61K0047-24 [I,C*]; A61K0047-24
                         [I,A]
                         A61K009/48H4
                  ECLA
                         A61K0009-00 [ICM]
 JP 53072814
                  IPCI
                         A61K0047-14 [I,C*]; A61K0047-14 [I,A]; A61K0009-48
                  TPCR
                         [I,C*]; A61K0009-48 [I,A]; A61K0047-06 [I,C*];
                         A61K0047-06 [I,A]; A61K0047-24 [I,C*]; A61K0047-24
                         [I,A]
                         A61K0047-00 [ICM]
 FR 2372635
                  IPCI
                         A61K0047-14 [I,C*]; A61K0047-14 [I,A]; A61K0009-48
                  IPCR
                         [I,C*]; A61K0009-48 [I,A]; A61K0047-06 [I,C*];
                         A61K0047-06 [I,A]; A61K0047-24 [I,C*]; A61K0047-24
                         [I,A]
     Capsules are prepared containing a drug solubilized by combination with a
AB
     spontaneously emulsifying oil. The oils contain a nonionic, surfactant as emulsifier. Phase diagrams for the formation of emulsions by a number of
     ternary oil-surfactant are presented. For example, a formulation containing
     diazepam 3.03, Miglyol 812 [37332-31-3] 29.09, Brij 35 [9002-92-0]
     48.49, and Span 80 [1338-43-8] 19.39 weight percent emulsified rapidly and
     spontaneously in water and could be dispensed into soft gelatin capsules.
ST
     solubilization drug oil emulsifier
IT
     Emulsions
        (of oils, pharmaceutical solubilization in relation to)
     Solubilization
IT
        (of pharmaceuticals, by spontaneously emulsifying oil)
     Pharmaceuticals
        (solubilization of, by spontaneously emulsifying oil)
IT
     Glycerides
     RL: BIOL (Biological study)
        (C8-12, pharmaceutical solubilization by spontaneously emulsifying
        formulation of)
                  9002-92-0
IT
     1338-43-8
     RL: BIOL (Biological study)
        (pharmaceutical solubilization by spontaneously emulsifying oil
        formulation containing)
```